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Docket No. GJE-6757C1

Serial No. 10/617,847

In the Specification

Please replace the following paragraph on page 3, lines 11-17 (2<sup>nd</sup> full paragraph):

For use in the invention, the active compound can be formulated in any suitable manner together with a conventional diluent or carrier. The active compound is preferably administered by the oral route; other suitable routes of administration include sublingual/buccal, transdermal, intramuscular, intranasal, rectal, parenteral, subcutaneous, pulmonary and topical. An effective dose of the active agent will depend on the nature and degree of the complaint, the age and condition of the patient and other factors known to those skilled in the art. A typical daily dosage may be 0.1 mg to 1g or 5 mg.

Clean Version of Amendment

Please replace the following paragraph on page 3, lines 11-17 (2<sup>nd</sup> full paragraph):

For use in the invention, the active compound can be formulated in any suitable manner together with a conventional diluent or carrier. The active compound is preferably administered by the oral route; other suitable routes of administration include sublingual/buccal, transdermal, intramuscular, intranasal, rectal, parenteral, subcutaneous, pulmonary and topical. An effective dose of the active agent will depend on the nature and degree of the complaint, the age and condition of the patient and other factors known to those skilled in the art. A typical daily dosage may be 0.1 mg to 1g or 5 g.